NEWBORN SCREENING CONFIRMED CASE REPORT PRIMARY PANEL OF 29 DISORDERS DOB 1/1/2008 – 12/31/2008		
Disorder	Abbreviation	Total
Congenital Hypothyroidism	CH	51
Congenital Adrenal Hyperplasia (21-hydroxylase deficiency)	CAH	5
Sickle Cell Anemia	Hb SS	8
Sickle Beta Thalassemia	Hb S/ßTh	0
Sickle C Disease	Hb S/C	1
Biotinidase Deficiency	BIOT	1
Galactosemia	GALT	2
Phenylketonuria	PKU	2
Maple Syrup Urine Disease	MSUD	0
Homocystinuria	HCY	1
Citrullinemia	CIT	1
Argininosuccinic Acidemia	ASA	0
Tyrosinemia Type 1	TYR I	0
Carnitine Uptake Defect	CUD	1
Medium-chain Acyl-CoA Dehydrogenase Deficiency	MCAD	4
Very Long-chain Acyl-CoA Dehydrogenase Deficiency	VLCAD	0
Long-chain 3-OH Acyl-CoA Dehydrogenase Deficiency	LCHAD	0
Trifunctional Protein Deficiency	TFP	0
Isovaleric Acidemia	IVA	0
Glutaric Acidemia Type 1	GA I	0
Hydroxymethylglutaric CoA Lyase Deficiency	HMG	0
3-Methycrotonyl CoA Carboxylase Deficiency	3MCC	1
Multiple Carboxylase Deficiency	MCD	0
Methylmalonic Acidemia (mutase deficiency)	MUT	0
Methylmalonic Acidemia (Cbl A,B)	Cbl A,B	0
Propionic Acidemia	PROP	0
Beta-Ketothiolase Deficiency	BKT	0
Cystic Fibrosis	CF	14
Hearing Loss	HEAR	122
TOTAL (92 bloodspot, 122 hearing)		**214**

As of 4-23-2009